

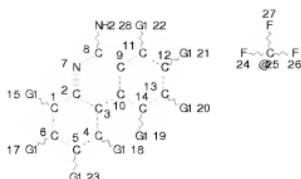
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=> d hi s
(FI LE 'HOME' ENTERED AT 12:31:33 ON 12 JAN 2010)
  FI LE 'LREG STR' ENTERED AT 12:31:51 ON 12 JAN 2010
L1    STR
  FI LE 'REG STR' ENTERED AT 12:32:38 ON 12 JAN 2010
L2    50 S L1
L3    7168 S L1 FUL
L4    1819 S L3 AND NRS=1
L5    712 S L4 AND X ELS
  FI LE 'LREG STR' ENTERED AT 12:33:43 ON 12 JAN 2010
L6    STR L1
  FI LE 'REG STR' ENTERED AT 12:48:12 ON 12 JAN 2010
L7    26 S L6 SSS SAM SUB=L3
L8    STR L6
L9    3 S L8 SSS SAM SUB=L3
L10   73 S L8 SSS FUL SUB=L3
L11   42 S L10 AND NG=1
  FI LE 'CAPLUS' ENTERED AT 12:55:15 ON 12 JAN 2010
L12   1178 S L1
L13   29 S L12(L) THU RL
L14   11 S L13 AND PY=2004
  FI LE 'LREG STR' ENTERED AT 13:07:46 ON 12 JAN 2010
L15   STR L8
  FI LE 'REG STR' ENTERED AT 13:08:21 ON 12 JAN 2010
L16   1 S L15 SSS FUL SUB=L3
  FI LE 'CAPLUS' ENTERED AT 13:08:47 ON 12 JAN 2010
L17   57838 S L1 DMG
L18   57838 S L1(5A) PHARMACEUTICAL?
L19   157 S L18(5A) PHARMACEUTICAL?
L20   86 S L19 AND PY=2003
L21   3 S L19(5A) CARRIER
  FI LE 'STNGUI DE' ENTERED AT 13:17:48 ON 12 JAN 2010
  FI LE 'LREG STR' ENTERED AT 13:41:39 ON 12 JAN 2010
L22   STR L15
  FI LE 'REG STR' ENTERED AT 13:42:27 ON 12 JAN 2010
L23   15 S L22 SSS FUL SUB=L3
  FI LE 'CAPLUS' ENTERED AT 13:43:11 ON 12 JAN 2010
L24   37 S L23
L25   6 S L24(L) THU RL

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GRAPH ATTRIBUTES:
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L26 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2010 ACS on STN
AN 2004054785 CAPLUS

DN 140 121957 TI Isolation of drugs active against mammalian prions using a yeast-based screen in assay

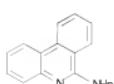
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OS Station Biologique, Centre Cyclotron Laboratory, C.N.R.S., Bretagne, 29660, FR
SO Nature Biotechnology (2003), 21(9), 1076-1081

PB Nature Publishing Group
DT Journal
LA English

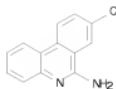
AB We have developed a rapid, yeast-based, two-step assay to screen for anti-prion drugs. The method allowed us to identify several compounds effective against budding yeast prions responsible for the [PSI⁺] and [URE3] phenotypes. These inhibitors include the kastellipadiines, a new class of compounds, and two previously known ones, 6-phenanthridine and 6-chlorophenanthridine. A second pool of compounds, which share structural similarities with the kastellipadiines, were also active in the assay. The compounds isolated here were also active in promoting mammalian prion clearance. These results validate the present method as an efficient high-throughput screening approach to identify new prion inhibitors and further reinforce the suggestion that biochemical pathways controlling prion formation and maintenance are conserved from yeast to humans.

IT 651055-79-7
RN 651055-79-7
RL PAC (Pharmacological activity); THU (Therapeutic use); BICL (Biological study); USES (Uses)
(Isolation of drugs active against mammalian prions using a yeast-based screen in assay)

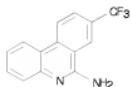
RN 832-66-8 CAPLUS
CN 6-Phenanthridinamine (CA INDEX NAME)



RN 651055-79-7 CAPLUS
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RN 651055-83-3 CAPLUS
CN 6-Phenanthridinamine, 8-(trifluoromethyl)- (CA INDEX NAME)



OSC G 66 THERE ARE 66 CAPLUS RECORDS THAT CITE THIS RECORD (66 CITINGS)
RE CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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